CLAIMS

An amide derivative represented by general formula
 below or a salt thereof.

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$$Z$$
 N
 X
 R^3
 (I)

(In the formula, the symbols represent the following meanings:

Z: 1,2,4-oxadiazol-3-yl, 4-oxazolyl, 1,2,3-triazol-210 yl or 2-pyridyl group,

A: an aryl which may have a substituent(s),
heteroaryl which may have a substituent(s), saturated
hydrocarbon ring-fused aryl which may have a substituent(s)
or saturated heterocyclic ring-fused aryl group which may
have a substituent(s), provided that the saturated
hydrocarbon ring-fused aryl or saturated heterocyclic ringfused aryl group is bonded to a nitrogen atom via a carbon
atom in an aromatic ring,

X: CO or SO_2 ,

R³: an alkyl which may have a substituent(s), alkenyl which may have a substituent(s), alkynyl which may have a substituent(s), cycloalkyl which may have a substituent(s), cycloalkenyl which may have a substituent(s), aryl which

may have a substituent(s), or heterocyclic group which may
have a substituent(s) or NRaRb,

Ra and Rb: which are the same or different from each other, H, a lower alkyl, lower alkenyl, lower alkynyl, cycloalkyl, cycloalkenyl, aryl, 5- or 6-membered monocyclic heteroaryl which has 1 to 4 hetero atoms selected from a group consisting of N, S and O, or lower alkylene-aryl group.

- The amide derivative or a salt thereof according to
 Claim 1, wherein X is CO.
 - 3. The amide derivative or a salt thereof according to Claim 1, wherein A is an aryl group selected from a phenyl and naphthyl group; a heteroaryl group selected from a pyridyl, pyrimidinyl, benzofuranyl, benzothienyl,
- benzothiadiazolyl, benzothiazolyl, benzoxazolyl,
 benzoxadiazolyl, benzimidazolyl, indolyl, isoindolyl,
 indazolyl, imidazopyridyl and indolidinyl group; a
 saturated hydrocarbon ring-fused aryl group selected from
 4-indanyl, 5-indanyl, 5,6,7,8-tetrahydronaphthalene-1-yl
- and 5,6,7,8-tetrahydronaphthalene-2-yl; or a saturated heterocyclic ring-fused aryl group selected from a 3,4-dihydro-2H-1,4-benzoxadinyl, 3,4-dihydro-2H-1,4-benzothiadinyl, 1,3-benzodioxolyl, 2,3-dihydro-1,4-benzodioxynyl, chromanyl, isochromanyl, 3,4-dihydro-2H-1-
- benzothiopyranyl, 3,4-dihydro-1H-2-benzothiopyranyl,
 indolinyl, isoindolinyl, 1,2,3,4-tetrahydroquinolyl, and

- 1,2,3,4-tetrahydroisoquinolyl group; the aryl, heteroaryl, saturated hydrocarbon ring-fused aryl and saturated heterocyclic ring-fused aryl each may have 1 to 5 substituents selected from Group D1:
- R³ is a cycloalkyl selected from cyclopentyl, cyclohexyl and cycloheptyl, cycloalkenyl selected from cyclopentenyl and cyclohexenyl, aryl selected from phenyl and naphthyl, saturated heterocyclic ring-fused aryl selected from 1,3-benzodioxolyl, 2,3-dihydro-1,4-benzodioxinyl, 3,4-dihydro-
- 2H-1-benzothiopyranyl and 3,4-dihydro-1H-2-benzothiopyranyl, heteroaryl selected from pyridyl, pyrimidinyl, benzofuranyl, benzothienyl, benzothiadiazolyl, benzothiazolyl, benzoxazolyl, benzoxadiazolyl, benzimidazolyl, indolyl, isoindolyl, indazolyl,
- imidazopyridyl and indolidinyl group, or 5- to 8-membered saturated heterocyclic group selected from tetrahydro-2H-pyranyl, tetrahydro-2H-thiopyranyl, thiepanyl, thiocanyl, thiabicyclo[3.1.0]hexanyl, perhydro-1,3-thiazinyl, pyrrolidinyl, imidazolidinyl, pyrazolidinyl, piperadinyl,
- azepanyl, diazepanyl, piperidinyl, morpholinyl and thiomorpholinyl group, the cycloalkyl, cycloalkenyl, aryl, saturated heterocyclic ring-fused aryl, heteroaryl and 5-to 8-membered saturated heterocyclic group each may have 1 to 5 substituents selected from Group D1 and the sulfur
- 25 atom of the ring may form oxide or dioxide; and

Group D1: lower alkyl, phenyl, halogeno lower alkyl, COOH, COO-lower alkyl, CO-lower alkyl, halogen atoms, NO2, CN, OH, lower alkylene-OH, lower alkylene-O-lower alkyl, Olower alkyl, O-halogeno lower alkyl, O-lower alkylene-OH, O-lower alkylene-O-lower alkyl, O-lower alkylene-COOH, Olower alkylene-COO-lower alkyl, O-lower alkylene-NH2, Olower alkylene-NH-lower alkyl, O-lower alkylene-N(lower alkyl)2, O-lower alkylene-(a nitrogen-containing saturated heterocyclic group which may be substituted with a lower 10 alkyl group(s)), O-phenyl, O-lower alkylene-phenyl, NH2, NH-lower alkyl, NH-lower alkylene-OH, NH-lower alkylene-Olower alkyl, NH-lower alkylene-NH2, NH-lower alkylene-NHlower alkyl, NH-lower alkylene-N(lower alkyl)2, NH-lower alkylene-(a nitrogen-containing saturated heterocyclic group which may be substituted with a lower alkyl 15 group(s)), N(lower alkyl)₂, (a nitrogen-containing saturated heterocyclic group which may have a substituent(s) selected from lower alkyl and lower alkylene-COORa), NHCO-lower alkyl, N(lower alkyl)CO-lower 20 alkyl, CONH₂, CONH-lower alkyl, CON(lower alkyl)₂, =O(oxo), SH, S-lower alkyl, SO-lower alkyl, and SO₂-lower alkyl.

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- 4. The amide derivative or a salt thereof according to Claim 3, wherein A is a group selected from a phenyl,
- pyridyl, benzothiazolyl, indazolyl, 5-indanyl, 1,3benzodioxolyl and indolinyl group, all of which may have 1

- to 3 substituents selected from a group consisting of a lower alkyl, lower alkylene-O-lower alkyl, CF₃, halogen atoms, CO-lower alkyl, OH, O-lower alkyl, CN, OCF₃, O-lower alkylene-OH, O-lower alkylene-O-lower alkyl, NH₂, NH-lower alkyl, N(lower alkyl)₂, NH-lower alkylene-OH, NH-lower alkylene-O-lower alkyl and O-lower alkylene-phenyl; and R³ is a group selected from a cyclohexyl, phenyl, naphthyl, pyridyl, pyrimidinyl, benzothiazolyl, benzooxadiazolyl, thiabicyclo[3.1.0]hexanyl, tetrahydro-2H-pyranyl,
- thiomorpholinyl, tetrahydro-2H-thiopyranyl and perhydro-1,3-thiazinyl group, all of which may be substituted with 1 or 2 substituents selected from halogen atoms, CN, =0, OH, O-lower alkyl, lower alkylene-OH and CONH₂ and the sulfur atom of the ring may form oxide or dioxide.
- 15 5. The amide derivative or a salt thereof according to Claim 1, wherein Z is 1,2,4-oxadiazol-3-yl group.
 - 6. The amide derivative or a salt thereof according to Claim 1, wherein Z is 4-oxazolyl group.
- 7. The amide derivative or a salt thereof according to Claim 1, wherein A is a group selected from a phenyl and 5-indanyl group, all of which may have 1 to 4 substituents selected from a group consisting of a lower alkyl, O-lower alkyl and halogen atoms; X is CO; and R³ is 1,1-dioxidotetrahydro-2H-thiopyran-4-yl.
- 25 8. The amide derivative or a salt thereof according to Claim 7, wherein A is a phenyl, which is substituted a

methyl group and may further have 1 or 2 substituents selected from a group consisting of methyl and halogen atoms.

- 9. The amide derivative or a salt thereof according to Claim 7, wherein A is 5-indanyl group.
- 10. The amide derivative according to Claim 1, selected from N-(2,6-dimethylphenyl)-N-(2-{[4-(1,3-oxazol-4-yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-thiopyran-4-carboxamide 1,1-dioxide; N-(4-methylphenyl)-N-(2-{[4-(1,3-oxazol-4-yl)phenyl)-N-(2-(1,3-oxazol-4-yl)phenyl)-N-(2-(1,3-oxazol-4-yl)phenyl)-N-(2-(1,3-oxazol-4-yl)phenyl)-N-(2-(1,3-oxazol-4-yl)phenyl)-N-(2-(1,3-oxazol-4-yl)phenyl)-N-(2-(1,3-oxazol-4-yl)phenyl)-N-(2-(1,3-oxazol-4-yl)phenyl)
- oxazol-4-yl)phenyl]amino}-2-oxoethyl)tetrahydro-2Hthiopyran-4-carboxamide 1,1-dioxide; N-(3-methylphenyl)-N(2-{[4-(1,3-oxazol-4-yl)phenyl]amino}-2
 - oxoethyl)tetrahydro-2H-thiopyran-4-carboxamide 1,1-dioxide; $N-(2-methylphenyl)-N-(2-\{[4-(1,3-oxazol-4-yl)phenyl]amino\}-$
- 2-oxoethyl)tetrahydro-2H-thiopyran-4-carboxamide 1,1-dioxide; N-(2,4-dimethylphenyl)-N-(2-{[4-(1,3-oxazol-4-yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-thiopyran-4-carboxamide 1,1-dioxide; N-(3,4-dimethylphenyl)-N-(2-{[4-(1,3-oxazol-4-yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-
- thiopyran-4-carboxamide 1,1-dioxide; N-(2,3-dihydro-1Hinden-5-yl)-N-(2-{[4-(1,3-oxazol-4-yl)phenyl]amino}-2oxoethyl)tetrahydro-2H-thiopyran-4-carboxamide 1,1-dioxide;
 N-(4-chloro-3-methylphenyl)-N-(2-{[4-(1,3-oxazol-4yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-thiopyran-4-
- 25 carboxamide 1,1-dioxide; N-(3-fluoro-4-methylphenyl)-N-(2-{[4-(1,3-oxazol-4-yl)phenyl]amino}-2-oxoethyl)tetrahydro-

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2H-thiopyran-4-carboxamide 1,1-dioxide; N-(3-fluoro-2,4-
    dimethylphenyl)-N-(2-{[4-(1,3-oxazol-4-yl)phenyl]amino}-2-
    oxoethyl)tetrahydro-2H-thiopyran-4-carboxamide 1,1-dioxide;
    N-(3,5-difluoro-4-methylphenyl)-N-(2-{[4-(1,3-oxazol-4-
    yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-thiopyran-4-
    carboxamide 1,1-dioxide; N-(2-fluoro-4-methylphenyl)-N-(2-
    {[4-(1,3-oxazol-4-yl)phenyl]amino}-2-oxoethyl)tetrahydro-
    2H-thiopyran-4-carboxamide 1,1-dioxide; N-(2,3-
    dimethylphenyl)-N-(2-\{[4-(1,2,4-oxadiazol-3-
10
    yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-thiopyran-4-
    carboxamide 1,1-dioxide; N-(2,4-dimethylphenyl)-N-(2-{[4-
    (1,2,4-oxadiazol-3-yl)phenyl]amino}-2-oxoethyl)tetrahydro-
    2H-thiopyran-4-carboxamide 1,1-dioxide; N-(2,6-
    dimethylphenyl)-N-(2-\{[4-(1,2,4-oxadiazol-3-
15
    yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-thiopyran-4-
    carboxamide 1,1-dioxide; N-(4-fluoro-2,6-dimethylphenyl)-N-
    (2-\{[4-(1,2,4-oxadiazol-3-yl)phenyl]amino\}-2-
    oxoethyl)tetrahydro-2H-thiopyran-4-carboxamide 1,1-dioxide;
    N-(2,3-dihydro-1H-inden-5-y1)-N-(2-{[4-(1,2,4-oxadiazol-3-
20
    yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-thiopyran-4-
    carboxamide 1,1-dioxide; N-(3-fluoro-4-methylphenyl)-N-(2-
    {[4-(1,2,4-oxadiazol-3-yl)phenyl]amino}-2-
    oxoethyl)tetrahydro-2H-thiopyran-4-carboxamide 1,1-dioxide;
    N-(4-chloro-3-methylphenyl)-N-(2-{[4-(1,2,4-oxadiazol-3-
25
    yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-thiopyran-4-
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carboxamide 1,1-dioxide; and N-(3-fluoro-2,4-

dimethylphenyl)-N-(2-{[4-(1,2,4-oxadiazol-3yl)phenyl]amino}-2-oxoethyl)tetrahydro-2H-thiopyran-4carboxamide 1,1-dioxide.

- 11. A pharmaceutical composition which comprises the amide derivative or a salt thereof according to Claim 1 and a pharmaceutically acceptable carrier.
 - 12. The pharmaceutical composition according to claim 11 which is an anti-herpesvirus drug.
- 13. A method for treating diseases in which herpesvirus

 10 is involved which comprises administering to a patient in

 11 need of such treatment a therapeutically effective amount

 12 of an amide derivative or a salt thereof according to Claim

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